

REPORTING CHANGE

NOTIFICATION DATE: December 6, 2012 **EFFECTIVE DATE:** January 2, 2013

THINPREP DIAGNOSTIC WITH HUMAN PAPILLOMAVIRUS (HPV) REFLEX

Test ID: 83343 Secondary ID: 83343

EXPLANATION: Effective January 2, 2013, there will be a slight modification in the diagnostic terms used in reporting Pap tests. Cases with a diagnosis of Low Grade or High Grade Squamous Intraepithelial Lesion (LSIL or HSIL respectively), will be reported with only the appropriate Bethesda terminology. Consequently, the corresponding CIN/dysplasia terminology will no longer be used in conjunction with these diagnoses.

CURRENT REPORTING TERMINOLOGY:

Low grade squamous intraepithelial lesion. Consistent with mild dysplasia (CIN 1) Low grade squamous intraepithelial lesion. Consistent with mild dysplasia with associated HPV Changes (CIN 1)

High grade squamous intraepithelial lesion. Consistent with moderate dysplasia (CIN 2) High grade squamous intraepithelial lesion. Consistent with severe dysplasia (CIN 3) High grade squamous intraepithelial lesion. Consistent with carcinoma in situ (CIN 3)

NEW REPORTING TERMINOLOGY:

Low grade squamous intraepithelial lesion.

High grade squamous intraepithelial lesion.

QUESTIONS: Contact your Mayo Medical Laboratories' Regional Manager Richard R. Einerson, Mayo Medical Laboratories' Technical Support Telephone: 800-533-1710